**Amendments to the Claims:** 

This listing of claims will replace all prior versions, and listings, of claims in the

application:

**Listing of Claims:** 

Claim 1 (previously presented): The insulin regulator construct of Claim 9, wherein the

nucleotide sequence comprises:

a) a glucose response element (GIRE) of a liver-pyruvate (L-PK)

gene promoter; and

b) an insulin-sensitive element of an insulin-like growth factor binding

protein-1 (IGFBP-1) basal promoter.

Claim 2 (original): The insulin regulator construct of Claim 1, wherein:

said glucose response element comprises a hepatic nuclear

factor-4 (HNF-4) binding site and a glucose responsive site.

Claim 3 (original): The insulin regulator construct of Claim 2, further comprising:

a plurality of said glucose response elements.

Page 3 of 17

Appl. No.: 09/972,916

Amdt. dated September 1, 2009

Reply to Office Action of April 2, 2009

Claim 4 (original): The insulin regulator construct of Claim 2, wherein:

the sequence of said HNF-4 binding site and said glucose responsive site is in a native orientation.

Claim 5 (original): The insulin regulator construct of Claim 2, wherein:

the sequence of said HNF-4 binding site and said glucose responsive site is reversed from a native orientation.

Claim 6 (original): The insulin regulator construct of Claim 1, wherein:

said glucose response element is inserted upstream of said insulin-sensitive element in an insulin-like growth factor binding protein-1 (IGFBP-1) basal promoter.

Claim 7 (original): The insulin regulator construct of Claim 1, wherein:

said glucose response element comprises a nucleotide sequence set forth in SEQ ID NO.: 1.

Claim 8 (original): The insulin regulator construct of Claim 1, wherein:

said insulin-sensitive element comprises a nucleotide sequence set forth in SEQ ID NO.: 2.

Claim 9 (previously presented): An insulin regulator construct, comprising:

a) a nucleotide sequence set forth in one of SEQ ID NO.: 3, SEQ ID

NO.: 4, SEQ ID NO.: 5, and SEQ ID NO.: 6; and

b) a sequence encoding insulin or proinsulin operably linked to the

promoter element of said construct.

Claim 10 (previously presented): The insulin regulator construct of Claim 9, which is

not stimulated by exposure to lactate or fructose.

Claim 11 (previously presented): The insulin regulator construct of Claim 9, which is

stimulated by exposure to glucose and inhibited by exposure to insulin.

Claim 12 (previously presented): A vector comprising the construct of Claim 9.

Claim 13 (previously presented): An adenoviral vector comprising the construct of

Claim 9.

Claim 14 (previously presented): The construct of Claim 9, wherein said construct

comprises a transgene.

Claim 15 (previously presented): A pharmaceutical composition comprising the

construct of Claim 9 and a pharmaceutically acceptable carrier or diluent.

Claim 16 (canceled).

Claim 17 (withdrawn - currently amended): A method of treating or preventing diabetic

conditions in a subject by administering an effective amount of the construct of Claim

[[1]] <u>9</u>.

Claim 18 (withdrawn - currently amended): A method of regulating insulin production in

a subject by administering an effective amount of the construct of Claim [[1]] 9.

Claim 19 (withdrawn - currently amended): A method of modulating hyperglycemia,

while avoiding severe hypoglycemia, in a subject by administering an effective amount

of the construct of Claim [[1]] 9.

Claim 20 (withdrawn - currently amended): A method of increasing fat catabolism in a

subject by administering an effective amount of the construct of Claim [[1]] 9.

Claim 21 (withdrawn - currently amended): A method of reducing protein catabolism in

a subject by administering an effective amount of the construct of Claim [[1]] 9.

Page 6 of 17